IN THE CLAIMS

- 1. (Previously Presented) An isolated polynucleotide; which encodes a protein comprising the amino acid sequence of SEQ ID NO: 2.
 - 2. (Cancelled)
 - 3. (Original) A vector comprising the isolated polynucleotide of Claim 1.
 - 4. (Original) A host cell comprising the isolated polynucleotide of Claim 1.
 - 5. (Previously Presented) The host cell of Claim 4, which is a Corynebacterium.
- 6. (Previously Presented) The host cell of Claim 4, wherein said host cell is selected from the group consisting of Corynebacterium glutamicum, Corynebacterium acetoacidophilum, Corynebacterium melassecola, Corynebacterium thermoaminogenes, and Brevibacterium flavum.

7.-9. (Cancelled)

- 10. (Previously Presented) A method for making an OxyR transcriptional regulator protein, comprising:
- a) culturing the host cell of Claim 4 for a duration of time under conditions suitable for expression of an OxyR transcriptional regulator protein; and
 - b) collecting the OxyR transcriptional regulator protein.

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- 11. (Previously Presented) An isolated polynucleotide; which comprises nucleotides491 to 1471 of SEQ ID NO: 1.
- 12. (Previously Presented) An isolated polynucleotide, which is fully complementary to nucleotides 491 to 1471 of SEQ ID NO: 1.
 - 13.-18. (Cancelled)
 - 19. (Original) A vector comprising the isolated polynucleotide of Claim 11.
 - 20. (Original) A host cell comprising the isolated polynucleotide of Claim 11.
 - 21. (Previously Presented) The host cell of Claim 20, which is a Corynebacterium.
- 22. (Previously Presented) The host cell of Claim 20, wherein said host cell is selected from the group consisting of Corynebacterium glutamicum, Corynebacterium acetoacidophilum, Corynebacterium melassecola, Corynebacterium thermoaminogenes, and Brevibacterium flavum.
 - 23.-25. (Cancelled)
- 26. (Previously Presented) A method for making an OxyR transcriptional regulator protein, comprising:

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- a) culturing the host cell of Claim 20 for a duration of time under conditions suitable for expression of an OxyR transcriptional regulator protein; and
 - b) collecting the OxyR transcriptional regulator protein.
 - 27.-28. (Cancelled)
 - 29. (Original) Corynebacterium glutamicum DSM 13457.
 - 30.-39. (Cancelled)
- 40. (Previously Presented) A method for making an L-amino acid comprising: culturing in a suitable medium a cell comprising a polynucleotide encoding SEQ ID NO:2, and

recovering the L-amino acid,

wherein said cell overexpresses said polynucleotide and wherein said overexpression is achieved by increasing the copy number of said polynucleotide or operably linking to said polynucleotide a promoter or expression cassette to increase the expression of said polynucleotide.

- 41. (Previously Presented) The method of Claim 40, wherein said L-amino acid is L-lysine.
- 42. (Previously Presented) The method of Claim 40, wherein said cell is a Corynebacterium.

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43. (Previously Presented) The method of Claim 40, wherein said cell is selected from the group consisting of Corynebacterium glutamicum, Corynebacterium acetoacidophilum, Corynebacterium melassecola, Corynebacterium thermoaminogenes, and Brevibacterium flavum.

44.-49. (Cancelled)

- 50. (Currently Amended) A modified <u>Corynebacterium</u> Cornynebacterium comprising multiple copies of the polynucleotide of Claim 1.
- 51. (Previously Presented) A modified *Corynebacterium* comprising multiple copies of the polynucleotide of Claim 11.
- 52. (Previously Presented) A *Corynebacterium* modified to contain a polynucleotide encoding SEQ ID NO:2 under the control of an exogenous promoter or expression cassette, wherein the expression of the gene product of said polynucleotide is increased relative to a corresponding, unmodified *Corynebacterium*.
 - 53. The isolated polynucleotide of Claim 1 which comprises SEQ ID NO: 1.